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PATENT AND TRADEMARK OFFICEATTORNEY DOCKET NO.  
P107424-00001TRANSMITTAL LETTER TO THE UNITED STATES  
DESIGNATED/ELECTED OFFICE (DO/EO/US)  
CONCERNING A FILING UNDER 35 U.S.C. 371

DATE: March 27, 2000

U.S. APPLN. NO.  
(IF KNOWN, SEE 37 CFR 1.5)

09 / 508519

INTERNATIONAL APPLICATION NO. PCT/JP98/04286	INTERNATIONAL FILING DATE September 25, 1998	PRIORITY DATE CLAIMED September 27, 1997
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TITLE OF INVENTION: COATING COMPOSITION AND OPTICAL RECORDING MEDIUM

APPLICANT(S) FOR DO/EO/US: Tsuyoshi KOMAKI

1.  This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.  
(THE BASIC FILING FEE IS ATTACHED)
  2.  This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
  3.  This express request to begin national examination procedures (35 U.S.C. 371(f) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT articles 22 and 39(1).
  4.  A proper demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
  5.  A copy of the International Application as filed (35 U.S.C. 371(c)(2))
    - a.  is transmitted herewith (required only if not transmitted by the International Bureau).
    - b.  has been transmitted by the International Bureau.
    - c.  is not required, as the application was filed in the United States Receiving Office (RO/US)
  6.  A translation of the International Application into English (35 U.S.C. 371(c)(2)).
  7.  Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
    - a.  are transmitted herewith (required only if not transmitted by the International Bureau).
    - b.  have been transmitted by the International Bureau.
    - c.  have not been made; however, the time limit for making such amendments has NOT expired.
    - d.  have not been made and will not be made.
  8.  A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
  9.  An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
  10.  A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).
- Items 11. to 16. below concern other document(s) or information included:
11.  An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
  12.  An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
  13.  A **FIRST** preliminary amendment.  
 A **SECOND** or **SUBSEQUENT** preliminary amendment.
  14.  A substitute specification.
  15.  A change of power of attorney and/or address letter.
  16.  Other items or information:  
Check No. 288046  
Drawing(s) 1 sheet  
International Search Report  
International Preliminary Examination Report (PCT/IPEA/409)  
Front Page of International Bulletin No. WO99/16835

416 Rec'd PCT/PTO 27 MAR 2000

U.S. APPLN. NO. (IF KNOWN, SEE 37 C.F.R. 1.50) <b>09/508519</b>		INTERNATIONAL APPLICATION NO. PCT/JP98/04286		ATTORNEY DOCKET NO. P107424-00001
				DATE: March 27, 2000
<p>17. <input checked="" type="checkbox"/> The following fees are submitted:</p> <p><b>Basic National Fee (37 CFR 1.492(a)(1)-(5):</b></p> <p>Search Report has been prepared by the EPO or JPO.....\$840.00      International preliminary examination fee paid to USPTO (37 CFR 1.482)....\$670.00      No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)).....\$760.00      Neither international preliminary examination fee (37 CFR 1.482) or international search fee (37 CFR 1.445(a)(2)) paid to USPTO.....\$970.00      International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4) .....\$ 96.00</p>				CALCULATIONS   PTO USE ONLY
<b>ENTER APPROPRIATE BASIC FEE AMOUNT =</b>				\$ 840.00
Surcharge of \$130.00 for furnishing the oath or declaration later than <u>20</u> <u>30</u> months from the earliest claimed priority date (37 CFR 1.492(e)).				
Claims	Number Filed	Number Extra	Rate	
Total Claims	15 - 20 =	0	X \$ 18.00	
Independent Claims	1 - 3 =	0	X \$ 78.00	
Multiple dependent claim(s) (if applicable)			+\$260.00	260.00
<b>TOTAL OF ABOVE CALCULATIONS =</b>				\$ 1100.00
<input checked="" type="checkbox"/> Reduction by 1/2 for filing by small entity, if applicable. <input checked="" type="checkbox"/> Verified Small Entity statement must also be filed. <input checked="" type="checkbox"/> (Note 37 CFR 1.9, 1.27, 1.28).				
<b>SUBTOTAL =</b>				\$ 1100.00
Processing fee of \$130.00 for furnishing the English translation later the <u>20</u> <u>30</u> months from the earliest claimed priority date (37 CFR 1.492(f)). +				
<b>TOTAL NATIONAL FEE =</b>				\$ 1100.00
<input checked="" type="checkbox"/> Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +				\$ 40.00
<b>TOTAL FEES ENCLOSED =</b>				\$ 1140.00
				Amount to be refunded \$
				Charged \$
<p>a. <input checked="" type="checkbox"/> A check in the amount of \$ <u>1140.00</u> to cover the above fees is enclosed.</p> <p>b. <input type="checkbox"/> Please charge my Deposit Account No. <u>01-2300</u> in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.</p> <p>c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>01-2300</u>.</p>				
<p><b>NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.</b></p>				
SEND ALL CORRESPONDENCE TO:				
Arent Fox Kintner Plotkin & Kahn, PLLC 1050 Connecticut Avenue, N.W., Suite 600 Washington, D.C. 20036-5339 Telephone No. (202) 857-6000 Facsimile No. (202) 638-4810				
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09/508519

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416 Rec'd PCT/PTO 27 MAR 2000

## SPECIFICATION

COATING COMPOSITION AND OPTICAL RECORDING MEDIUM

## TECHNICAL FIELD

The present invention relates to a composition for coating which contains a radiation-curable compound, more particularly, a radiation-curable composition for coating which can form an ink receiving layer, on which image information and literature information can be printed with an ink jet method.

The present invention also relates to an optical recording medium having an ink receiving layer formed of the radiation-curable composition for coating.

## BACKGROUND ART

Recently, in an optical recording medium, there has been a greater demand for providing a surface opposite to a surface on which optical writing/reading is performed, with an ink receiving layer (label printing surface) and printing images or literatures representing a recorded content in the medium with an ink jet method so as to arrange the optical recording medium.

As an ink used for an ink jet recording method, there is usually used an aqueous one and, in many cases, a polyhydric alcohol or the like is added to this ink in order to prevent

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clogging of a nozzle.

On the other hand, as a substrate on which recording is made in the ink jet recording method, there have hitherto been used a normal paper, and a hydrophilized paper such as a specialty paper for ink jet which is provided with a porous ink receiving layer. As such an ink receiving layer, there are described, for example, the use of a water-soluble high polymer of poly (vinyl alcohol) and polyacrylic acid in Japanese Laid-open Patent Publication No.60-168651/1985, the use of hydroxyethyl cellulose in Japanese Laid-open Patent Publication No.60-262685/1985, the use of a mixture of carboxymethyl cellulose and polyethylene oxide in Japanese Laid-open Patent Publication No.61-181679/1986, and the use of a mixture of water-soluble cellulose and poly(vinylpyrrolidone) in Japanese Laid-open Patent Publication No.61-193879/1986. Since the water-soluble polymers described in these laid open publications are very hydrophilic, they have affinity with a hydrophilic substrate such as a paper and the like and, when an ink receiving layer composed of these water-soluble polymers is provided on the surface of a hydrophilic substrate, there arises no problem on adhesion between the substrate and the ink receiving layer.

However, for example, the water-soluble polymers described in the aforementioned laid open publications have little affinity with substrates having the surface of strong

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hydrophobicity such as an ultraviolet-curing resin and the like frequently used in a protecting membrane for the optical recording medium. For that reason, when an ink receiving layer composed of these water-soluble polymers is provided on the surface of a hydrophobic substrate, adhesion between the hydrophobic substrate and the ink receiving layer is worse and, after laminated, they are easily peeled and, additionally, since the ink receiving layer itself has the lower water-resistance, there are occurrence of an ink blot caused by a water droplet after printing and stickiness of the surface, which leads to a problem of practical use.

#### DISCLOSURE OF THE INVENTION

##### Object of the Invention

Therefore, an object of the present invention is to provide a composition for coating which overcomes the aforementioned problems of the prior art, and which can form, in one coating step, an ink receiving layer which is excellent in clearness of an image and quick-drying and which is excellent in adhesion with a hydrophobic substrate, the water-resistance and the stability of printing such as no occurrence of printing fall or no occurrence of separation of the whole layer even when wiped after wetting with water.

An another object of the present invention is to provide an optical recording medium having an ink receiving layer formed

of this composition for coating.

#### Summary of the Invention

The present inventor studied intensively and, as a result, found that the aforementioned objects can be attained by a composition comprising a radiation-curable acrylic compound with a water-absorbing organic filler and a water-absorbing inorganic filler added, which resulted in the completion of the present invention.

That is, the present invention is a composition for coating which comprises a radiation-curable compound (A) containing a monofunctional (meth) acrylic monomer and multifunctional more than bifunctional (meth) acrylic monomer, a water-absorbing organic filler (C) and a water-absorbing inorganic filler (D).

In the composition of the present invention, the radiation-curable compound (A) preferably contains 50-90% by weight of the monofunctional (meth) acrylic monomer and 10-50% by weight of the multifunctional more than bifunctional (meth) acrylic monomer.

In the composition of the present invention, the composition preferably contains 10-60% by weight of the water-absorbing organic filler (C) and 1-20% by weight of the water-absorbing inorganic filler (D) relative to the radiation-curable compound (A).

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In the present composition, a particle size of the filler (C) and (D) is preferably 0.001 to 20  $\mu\text{m}$ .

In the composition of the present invention, the composition preferably contains 0.001 to 10% by weight of a polymerization initiator relative to the radiation-curable compound (A).

The present invention is also a recording material having an ink receiving layer composed of a radiation-cured aforementioned composition for coating on at least one side of a hydrophobic supporting substrate.

The present invention is also an optical recording medium having an ink receiving layer composed of a radiation-cured aforementioned composition for coating on the surface opposite to the surface of the optical recording medium on which optical writing/reading is performed.

#### BRIEF DESCRIPTION OF THE DRAWING

Fig. 1 is a view showing a partial cross-section of one embodiment of an optical recording disk in accordance with the present invention.

#### DETAILED DESCRIPTION OF THE INVENTION

In the composition of the present invention, the radiation-curable compound (A) contains a monofunctional (meth) acrylic monomer and, as a cross-linking agent, a

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multifunctional more than bifunctional (meth) acrylic monomer.

The cured (meth) acrylic is highly hard, highly adhesive and highly cross-linked and has good adhesion to substrates having the hydrophobic surface such as ultraviolet-cured resins frequently used in a protecting film for optical recording medium.

In the present invention, examples of the monofunctional (meth) acrylic monomer, as a cross-linking agent, are not limited to specified ones but include alkyl (meth) acrylate such as ethyl (meth) acrylate, propyl (meth) acrylate, n-butyl (meth) acrylate, 2-ethylhexyl (meth) acrylate and the like, (meth) acrylic acid, (meth) acrylonitrile, (meth) acrylamide, N-substituted (meth) acrylamide, N-methylol (meth) acrylamide, hydroxyethyl (meth) acrylate, carboxyethyl (meth) acrylate, (meth) acryloylmorpholine, ethylene glycol mono (meth) acrylate, diethylene glycol mono (meth) acrylate, polyethylene glycol mono (meth) acrylate, dimethylaminopropyl (meth) acrylamide, tetrahydrofurfuryl (meth) acrylate and the like.

Among them, the high hydrophilic monomers having hydroxy group, carboxy group, amino group, morpholino group, or tetrahydrofurfuryl group are preferable.

Examples of the multifunctional more than bifunctional (meth) acrylic monomer as a cross-linking agent are not limited to specified ones and include ethylene glycol di (meth) acrylate, diethylene glycol di (meth) acrylate, polyethylene glycol di

(meth) acrylate, 1,6-hexane glycol di (meth) acrylate, pentaerythritol tetra (meth) acrylate, pentaerythritol (meth) acrylate, trimethylolpropane tri (meth) acrylate, trimethylolpropane di (meth) acrylate, and acrylic-modified urethane elastomer, as well as the aforementioned multifunctional (meth) acrylic monomers with a functional group such as COOH and the like introduced.

Among them, ethylene glycol di (meth) acrylate, diethylene glycol di (meth) acrylate, polyethylene glycol di (meth) acrylate, pentaerythritol tetra (meth)acrylate, pentaerythritol (meth) acrylate, trimethylolpropane tri (meth) acrylate, and trimethylolpropane di (meth) acrylate and the like are preferable.

In the composition of the present invention, it is preferable that the radiation-curable compound (A) contains 50-90% by weight of the monofunctional (meth) acrylic monomer and 10-50% by weight of the multifunctional more than bifunctional (meth) acrylic monomer.

When the multifunctional (meth) acrylic monomer, as a cross-linking agent, is less than 10% by weight, it tends to arise a problem of curability upon curing a composition or strength of a cured film. On the other hand, when the amount exceeds 50% by weight, there is a tendency that resulting cured film becomes too hydrophobic and printability is deteriorated.

It is more preferable that the radiation-curable compound

(A) contains 60-90% by weight of the monofunctional (meth) acrylic monomer and 10-40% by weight of the multifunctional (meth) acrylic monomer.

In addition, in the composition of the present invention, a (meth) acrylic oligomer or polymer, as a cross-linking agent, may be added therein as long as it does not deteriorate the objects of the present invention.

Since the radiation-curable compound (A) is polymerization-cured by a radiation, in particular, an ultraviolet-radiation, a photopolymerization initiator or a sensitizer is preferably incorporated into the present composition. The photopolymerization initiator or the sensitizer used is not limited to specified ones but may be selected from the conventional ones such as acetophenone type, benzoin type, benzophenone type and thioxanthone type compounds. As the photopolymerization initiator or the sensitizer, a plurality of compounds may be used. The content of the photopolymerization initiator may be 0.001 to 10% by weight, preferably 0.5 to 9% by weight relative to the radiation-curable compound (A).

In the composition of the present invention, examples of the water-absorbing organic filler (C) are not limited to specified ones but include fillers having hydroxy group, carboxy group, amino group or the like on the surface and/or in the interior thereof, more particularly, animal protein

ultrafine particles such as Staphyloid AC-3832 (manufactured by Takeda Chemical Industries, Ltd.), Flowbeads EA209 (manufactured by Sumitomo Seika Chemical Co., Ltd.), collagen powder and the like.

In the composition of the present invention, examples of the water-absorbing inorganic filler (D) are not limited to specified ones but include high-purity alumina, silica gel, silica-alumina, talc, diatomaceous earth, calcium carbonate, calcium sulfate, synthetic zeolite and the like. More particularly, there are Silcron G-600 (manufactured by SCM Chemicals), Aerosil TT-600 (manufactured by Aerosil) and the like.

The aforementioned water-absorbing organic filler (C) has an action of enhancing printability after radiation-curation of the composition and, on the other hand, the aforementioned water-absorbing inorganic filler (D) has an action of enhancing coating property of the composition onto a substrate. Therefore, the composition of the present invention contains both the water-absorbing organic filler (C) and the water-absorbing inorganic filler (D) as an essential component. When the filler component is only the water-absorbing organic filler (C), there arises a problem of coating property and pot life and, when the filler component is only the water-absorbing inorganic filler (D), sufficient printability can not be obtained. Inclusion of both the organic

filler (C) and the inorganic filler (D) can afford excellent characteristics in coating property and printability.

It is preferred that the composition of the present invention contains 10-60% by weight of the water-absorbing organic filler (C) and 1-20% by weight of the water-absorbing inorganic filler (D) relative to the radiation-curable compound (A). When the amount of the organic filler (C) is less than 10% by weight, sufficient printability is not easily obtained and, on the other hand, the amount of the organic filler (C) exceeds 60% by weight, a cured film having sufficient hardness is not easily obtained. In addition, when the amount of the inorganic filler (D) is less than 1% by weight, coating property tends to be deteriorated and, on the other hand, when the amount of the inorganic filler (D) exceeds 20% by weight, coating property becomes to be deteriorated because of remarkable increase in viscosity. More preferable amount of the filler is 20-50% by weight of the water-absorbing organic filler (C) and 3 to 15% by weight of the water-absorbing inorganic filler (D) relative to the radiation-curable compound (A).

In addition, in the composition of the present invention, it is preferred that a particle size of the water-absorbing organic filler (C) and the water-absorbing inorganic filler (D) is 0.001 to 20  $\mu\text{m}$  respectively. When the particle size is less than 0.001  $\mu\text{m}$ , water-absorbing property tends to be deteriorated and, on the other hand, when the particle size

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exceeds 20  $\mu\text{m}$ , smoothness of the coated surface tends to be deteriorated and the film tends to be brittle. More preferable particle size of the water-absorbing organic filler (C) or the water-absorbing inorganic filler (D) is approximately 0.1 to 10  $\mu\text{m}$  respectively. Additionally, a particle size used in this specification is a value obtained by a particle size measuring method using a laser and, for example, can be measured using a microtrack manufactured by Nikkiso Co., Ltd. and the like. Additionally, the particle size measuring method using a laser is described, for example, in item "2.2 Interaction of particles and electromagnetic wave" (pp. 85-87) in Ultrafine Particles Handbook (published by Fuji Technosystem (K.K.) on September 5, 1990).

Furthermore, in the composition of the present invention, maleic acid, itaconic acid, N-vinylpyrrolidone and the like which can be copolymerized with an acrylic monomer may be added in addition to the aforementioned components as long as it does not deteriorate the objects of the present invention.

A recording sheet having an ink receiving layer suitable for an ink jet method is obtained by applying the present coating composition on one side or both sides of a hydrophobic supporting substrate such as polyester type resin, polycarbonate type resin, acrylic type resin and the like, and curing the composition by irradiating with radiation such as an ultraviolet ray.

For irradiating with radiation, an electron beam or the like may be irradiated instead of an ultraviolet ray and, in some cases, an ultraviolet ray may be irradiated after heating. The intensity of an ultraviolet ray to be irradiated to the coated film is usually  $50 \text{ mW/cm}^2$  or more, and the dose of irradiation is usually approximately  $200-2000 \text{ mJ/cm}^2$ . As an ultraviolet ray source, the conventional one such as a mercury lamp may be used. By an ultraviolet ray irradiation, the aforementioned respective compounds are radical-polymerized and cured.

An optical recording medium having an ink receiving layer suitable for an ink jet method is obtained by applying the coating composition of the present invention on a protecting film of the optical recording medium and curing the composition by irradiating with radiation such as an ultraviolet ray.

One example of the optical recording medium, an optical recording disk, has a layered structure shown in Fig. 1. That is, Fig. 1 is a view showing a partial cross-section of one example of an optical recording disk, wherein an optical recording disk (1) has a coated-type recording layer (3) having a dye as a main component, a reflection layer (5) and a protecting film (6) in this order on a substrate (2), and has, on the protecting film (6), an ink receiving layer (7) composed of the coating composition of the present invention which has been

radiation-cured.

The substrate (2) is formed of a resin or a glass which is substantially transparent (preferably, transmittance of 88% or more) to a recording light and a reproducing light (light having wavelength of around 600 to 900 nm, particularly semiconductor laser light having wavelength of around 770-800 nm, especially 780 nm). This makes possible to record on and reproduce from the back of a substrate.

The substrate (2) is a disk having normal size and, when used as a recordable CD, it has a thickness of around 1.2 mm and a diameter of 80 to 120 mm.

As a materiel for the substrate, it is preferred that resins are used. Various thermoplastic resins such as polycarbonate resin, acrylic resin, amorphous polyolefin, TPX and the like are suitable. The substrate (2) may be prepared according to the known method such as injection molding and the like. Upon this, it is preferred that the predetermined pattern such as a groove (21) or the like is formed on the substrate surface for tracking or addressing. In addition, a resin layer having the predetermined pattern such as a groove or the like may be formed according to a 2P method or the like after manufacturing the substrate (2).

The groove is preferably a spiral continuous-type groove and it is preferred that a depth thereof is 500 to 3000 Å, a width thereof is 0.2 to 1.1 μm, in particular 0.3 to 0.6 μm,

and a width of land (portion between neighboring grooves) is 0.5 to 1.4  $\mu\text{m}$ , in particular 1.0 to 1.3  $\mu\text{m}$ . By taking such a construction of the groove, a good tracking signal can be obtained without lowering the reflection level of the groove portion. In addition, the groove may be provided with concave and convex for an addressing signal. Furthermore, a recording light is preferably directed to a recording layer in the groove.

The recording layer (3) may contain only one kind of dye or may be in the state where two or more dyes are compatible.

When recording a CD signal, it is preferred that an attenuation coefficient of a recording light and reproducing light wavelength (imaginary part of complex refractive index)  $k$  of the recording layer (3) is 0.03 to 0.25. When  $k$  is less than 0.03, absorbing rate of the recording layer is decreased and it becomes difficult to perform recording with the conventional recording power. When  $k$  exceeds 0.25, the reflectivity is below 60% and it becomes difficult to perform reproduction with the CD specification. In addition, when  $k$  is 0.03 to 0.20, particularly 0.03 to 0.15, extremely preferable results are obtained.

In addition, it is preferred that refractive index (real part of complex refractive index)  $n$  is 1.8 to 4.0, more preferably 2.0 to 3.0. When  $n$  is < 1.8, the reflectivity tends to be decreased and a signal modulating degree becomes smaller, which makes difficult to produce with a CD player. In addition,

PCT/JP98/04286 10

in order to obtain  $n > 4.0$ , a raw material dye is obtained with difficulty.

Examples of the light-absorbing dye used in the recording layer (3) are not limited to specified ones as long as they have maximum absorption of 600 to 900 nm, preferably 600 to 800 nm, more preferably 650 to 750 nm and one or two or more may be selected from cyanine type, phthalocyanine type, naphthalocyanine type, anthraquinone type, azo type, triphenylmethane type, pyrylium or thiapyrylium salt type, squarylium type, croconium type, formazan type, metal complex type dye and the like depending upon the purpose.

As the cyanine type dye, cyanine dyes having an indolenine ring optionally having an aromatic condensed ring, particularly benzoindolenine ring are preferable. In addition, it is preferred that the indolenine rings are bonded via a pentamethylene chain.

In addition, it is preferred that the light-absorbing dye is used in admixture with a singlet oxygen quencher. Furthermore, it is also preferred that a dye cation and a singlet oxygen quencher anion which are ionically-bonded with each other are used as the light-absorbing dye.

As the quencher, metal complexes of acetylacetone type, bisdithiol type such as bisdithio- $\alpha$ -diketone type and bisphenyldithiol type, thiocatechol type, salicylaldehydeoxime type, thiobisphenolate type and the like

are preferable. In addition, amine type quenchers such as amine type compounds having a nitrogen radical cation, hindered amine and the like are suitable.

As a dye which constitutes the bonded system, cyanine dyes having an indolenine ring are preferable and, as the quencher, metal complex dye such as bisphenyldithiol metal complex and the like are preferable.

Further, although the quencher may be added independently from the light-absorbing dye or may be used in a bonded form, it is preferred that the quencher is added in an amount of 1 mole or less, particularly approximately 0.05 to 0.5 mole relative to 1 mole of total light-absorbing dyes. This can promote light-resistance.

As a dye used in the recording layer, those having n and k described above may be selected from the aforementioned light-absorbing dyes, dye-quencher mixtures, and dye-quencher bonded forms and, alternatively, the dyes may be synthesized by new molecular design.

In addition, k for a recording light and reproducing light of the dye varies in a range of around 0 to 2 depending upon the skeleton and substituent thereof and, thus, there is a limit of the skeleton and substituent upon selection of the dyes having k of 0.03 to 0.25. For that reason, in some cases, there arise a limit on coating solvents and impossible to coat depending upon the substrate materials. In addition, in some

PCT/JP98/04286

cases, vapor film forming can not be carried out. In addition, when new molecular design is carried out, there requires a great effort for design and synthesis.

This can be applied to the improvement of wavelength dependency. The wavelength of a semiconductor laser is usually in a range of  $\pm 10$  nm and, in a commercially available CD players, reflectivity of 70% or more need to be maintained at 770 nm to 790 nm. Generally, k value of the dyes has great wavelength dependency in many cases and, even when k value is suitable at 780 nm, k value becomes remarkably unsuitable at 770 nm or 790 nm in many cases. In such cases, suitable n and k values can be usually obtained in a range of  $780 \pm 10$  nm by mixing the second dye therein.

As a result, the limitation such as that of a coating solvent or the like is alleviated and it can make possible to use a dye which is easily synthesized and is low cost, a dye having good property, or a refractory dye.

When the recording layer (3) is a mixed dye layer, dyes to be used may be selected from those having  $n$  of 1.6 to 6.5 and  $k$  of 0 to 2.

In addition, upon measuring n and k, a recording layer is provided at a thickness of around 40 to 100 nm on the predetermined transparent substrate under the actual conditions to obtain a sample for measurement. Then, reflectivity through a substrate of this sample for measurement

or the reflectivity from a recording layer side is measured. The reflectivity is measured using a recording-reproducing light wavelength in a manner of mirror reflection (about 5°). In addition, transmittance of the sample is measured. From these measured values, n and k may be calculated according to, for example, Kyoritsuzensho "Optics" authored by Kozo Ishiguro pp. 168-178.

The recording layer may be formed by a spin coating method in which a coating solution containing a dye and an organic solvent is used and this coating solution is applied-spread onto a spinning substrate.

The organic solvent used in a coating solution for forming the recording layer may be conveniently selected from alcohol type, ketone type, ester type, ether type, aromatic type, and halogenated alkyl type solvents and the like depending upon a dye to be used. Organic solvents having two or more functional groups per molecule are suitable.

After spin coating, the formed film is dried as necessary.

The thickness of the recording layer thus formed is conveniently set depending upon the desired reflectivity and the like and is usually around 100 to 300 nm.

A reflection layer (5) is provided on the recording layer (3) in a manner of direct adhesion. The reflection layer (5) is composed of Au, Ag or Ag alloy, Au being preferable in view of high reflectivity and good anti-corrosion.

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As an Ag alloy, those containing one or more selected from Cu, Ti, V, Ta, Cr, Mo, W, Mn, Fe, Co, Rh, Ni, Pd, Pt, Au, Al and the like, in addition to Ag, are preferable. In order to obtain high reflectivity, it is preferred that the content of Ag is 40 atom % or more.

In order to form the reflection layer (5), various vapor film forming methods such as sputtering, deposition and the like may be used. In addition, it is preferred that the thickness of the reflection layer (5) is 20 nm or more. When the thickness is smaller, the reflectivity is liable to become insufficient.

By taking such a construction of the reflection layer, the reflectivity through the substrate when manufactured into an optical recording disk (unrecorded portion) is 60% or more, in particular 70% or more.

A protecting film (6) is provided on the reflection layer (5).

Although the protecting film (6) is preferably formed by a spin coating method in order to avoid a damage of the reflection layer (5), a screen printing method, a dipping method, a spray coating method and the like may be used. The conditions for forming the protecting film (6) are not particularly limited to specified ones but may be conveniently selected depending upon viscosity of the coating solution, desired thickness and the like.

The thickness of the protecting film is preferably around

1 to 20  $\mu\text{m}$ .

When the thickness is smaller, anti-corrosion effect of the reflection layer and the recording layer is liable to be insufficient. When the thickness is larger, cracks are liable to occur or warpages are liable to occur on a disk, deriving from shrinkage upon curing.

The protecting film is preferably composed of a radiation-cured resin. More particularly, it is preferably composed of a material obtained by radiation-curing a radiation-curable-type compound or its composition for polymerization. Examples thereof include monomers, oligomers and polymers in which groups cross-linkable or polymerizable by radiation are contained or introduced into a molecule, such as acrylic double bonds such as acrylic acid, methacrylic acid and ester compounds thereof, allyl double bonds such as diallyl phthalate, unsaturated double bonds such as maleic acid, and maleic derivatives which respond to an ionization energy and have unsaturated double bonds showing radical polymerizability. These are preferably multifunctional, in particular trifunctional or more and one kind of them may be used or two or more of them may be used together.

As a radiation-curable monomer, the compounds having a molecular weight of less than 2,000 are suitable and, as an oligomer, the compounds having a molecular weight of 2,000 to 10,000 are suitable. Examples thereof are styrene, ethyl

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acrylate, ethylene glycol diacrylate, ethylene glycol dimethacrylate, diethylene glycol diacrylate, diethyleneglycol methacrylate, 1,6-hexane glycol diacrylate, 1,6-hexane glycol dimethacrylate and the like. Particularly preferred are acryl group-containing monomers and/or oligomers such as pentaerythritol tetra (meth) acrylate, pentaerythritol (meth) acrylate, trimethylolpropane tri (meth) acrylate, trimethylolpropane di (meth) acrylate, acryl-modified urethane elastomer, or the foregoing with an introduced functional group such as COOH and the like, (meth) acrylate of phenol-ethylene oxide adduct, compounds in which a (meth) acrylate group or a  $\epsilon$ -caprolactone-acryl group is bonded to a pentaerythritol condensed ring shown in Japanese Laid-open Patent Publication No.1-165050/1989 (Japanese Examined Patent Publication No.8-27990/1996), special acrylates and the like. In addition to them, as a radiation-curable oligomer, there are acryl-modified oligoester acrylate and urethane elastomer, and the foregoing with an introduced group such as COOH and the like.

Alternatively, in addition to or instead of the aforementioned compounds, radiation-curable-type compounds obtained by radiation-responsively-modifying thermoplastic resin may be used. Examples of such a radiation-curable resin are resin in which groups cross-linkable or polymerizable by radiation are contained or introduced into a thermoplastic resin molecule, such as acrylic double bonds such as acrylic

acid, methacrylic acid or ester compounds thereof, allyl double bonds such as diallyl phthalate, and unsaturated bonds such as maleic acid, maleic derivatives which have unsaturated double bonds showing radical polymerizability. Examples of the thermoplastic resin which can be modified to radiation-curable resin are vinylchloride type copolymer, saturated polyester resin, poly (vinyl alcohol) type resin, epoxy type resin, phenoxy type resin, cellulose derivatives and the like. As other example of a resin which can be used for radiation-responsively-modifying, multifunctional polyester resin, polyether ester resin, poly (vinylpyrrolidone) resin and derivatives thereof (PVP olefin copolymer), polyamide resin, polyimide resin, phenol resin, spiroacetal resin, and acrylic type resins containing at least one of hydroxy group-containing acrylic ester and methacrylic ester as a polymerization component are also effective.

Since the coating composition for polymerization is cured by radiation, particularly ultraviolet radiation, it is preferred that a photopolymerization initiator or a sensitizer is contained in the composition for polymerization. The photopolymerization initiator or the sensitizer used is not limited to specified ones and examples thereof are as described above. The content of the photopolymerization initiator in the composition for polymerization is usually around 0.5 to 5% by weight. Such a composition for polymerization may be

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synthesized according to the conventional method or may be prepared using a commercially available compound.

As a composition containing a radiation-curable-type compound for forming the protecting film, a composition containing epoxy resin and an photo-cation polymerization catalyst is also suitably used.

As an epoxy resin, alicyclic epoxy resins are preferable and, particularly, those having two or more epoxy groups are preferable. As an alicyclic epoxy resin, one or more of 3,4-epoxycyclohexylmethyl-3,4-epoxycyclohexanecarboxylate, bis-(3,4-epoxycyclohexylmethyl)adipate, bis-(3,4-epoxycyclohexyl)adipate, 2-(3,4-epoxycyclohexyl-5,5-spiro-3,4-epoxy)cyclohexane-metha-dioxane, bis(2,3-epoxycyclopentyl)ether, vinylcyclohexenedioxide and the like are preferable. The epoxy equivalent of the alicyclic epoxy resin is not limited to specified ones but 60 to 300, particularly 100 to 200 is preferable in view of good curability.

As the photo-cation polymerization catalyst, any known ones may be used and particularly not limited to specified ones. For example, one or more of metal fluoroboric acid salt and boron trifluoride complex, bis (perfluoroalkyl-sulfonyl) methane metal salt, aryl diazonium compound, aromatic onium salt of 6A group element, aromatic onium salt of 5A group element, dicarbonylchelate of 3A-5A group element, thiopyrylium salt,

6A group element having MF<sub>6</sub> anion (provided that M is P, As or Sb), triarylsulfonium complex salt, aromatic iodonium complex salt, aromatic sulfonium complex salt and the like may be used. Particularly, it is preferred that one or more of polyarylsulfonium complex salt, aromatic sulfonium salt or iodonium salt of halogen-containing complex ion, and aromatic onium salt of 3A group element, 5A group element and 6A group element are used.

In addition, an photo-cation polymerization catalyst containing an organometallic compound and an organosilicon compound having photodegradability may be also used. Since such an photo-cation polymerization catalyst is non-strong acid type, adverse effects on the recording layer having high corrosiveness of a an optical magnetic recording disk can be avoided. As an organometallic compound, complex compounds in which alkoxy group, phenoxy group,  $\beta$ -diketonato group and the like are bonded to a metal atom such as Ti, V, Cr, Mn, Fe, Co, Ni, Cu, Zn, Al, Zr and the like are preferable. Among them, particularly, organoaluminum compound is preferable, more particularly, trismethoxyaluminium, trispropionatoaluminium, tristrifluoroacetylaluminium, and triethylacetacetatoaluminium are preferable.

The organosilicon compound having photodegradability produces a silanol by light irradiation such as ultraviolet ray or the like, and the silicon compounds having peroxy silano group,

*o*-nitrobenzyl group,  $\alpha$ -ketosilyl group and the like are preferable.

It is preferred that the content of the photo-cation polymerization catalyst in the composition is 0.05 to 0.7 part by weight, particularly 0.1 to 0.5 part by weight relative to 100 parts by weight of the epoxy resin.

Among them, those obtained by radiation-curing, particularly ultraviolet ray-curing a coated film containing the photopolymerization sensitizer or the initiator using a radiation-curable-type compound having an acryl group are preferable.

Upon formation of the protecting film, a coated film for the aforementioned composition for polymerization is irradiated with ultraviolet ray, electron beam or the like to cure it.

In addition, the protecting film may contain various pigment particles as necessary.

An ink receiving layer (7) composed of the coating composition of the present invention which has been radiation-cured is provided on the protecting film (6). The coating composition is applied, for example, by a screen printing method and this is irradiated with a radiation such as ultraviolet ray to cure, leading to an ink receiving layer suitable for an ink jet method.

## MODE FOR CARRYING OUT THE INVENTION

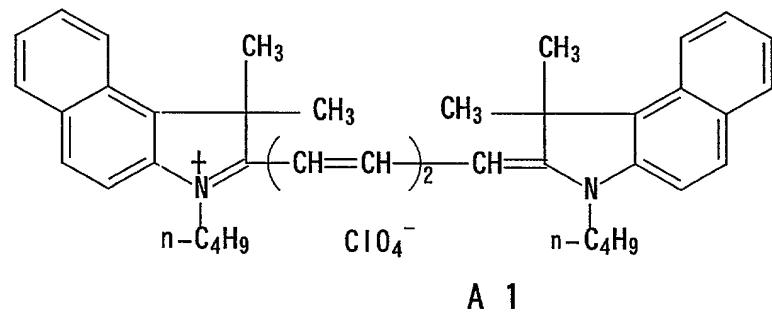
The present invention is illustrated by the following Examples but are not limited to them.

## [Example 1]

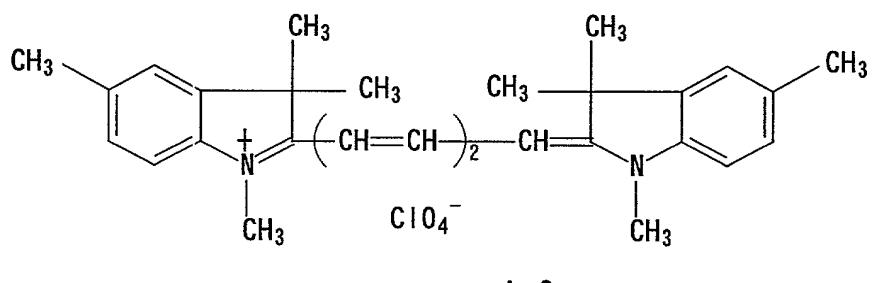
A Recordable-type compact disk (CD-R) having a construction shown in Fig. 1 was manufactured. First, up to formation of a protecting film (6) was performed. That is, a recording layer (3) containing a dye was formed on a polycarbonate resin substrate (2) having spiral continuous grooves, 120 mm in a diameter and 1.2 mm in a thickness with a spin coating method. A reflection layer (5) was formed on this recording layer (3) and a protecting film (6) was further formed thereon.

A coating solution used for forming the recording layer (3) contained a dye and an organic solvent and, as a dye, the following dyes A1 and A2 as well as singlet oxygen quencher Q were used, and the content of A1 in the dye was 60% by weight, that of A2 in the dye was 30% by weight and that of the singlet oxygen quencher was 10% by weight.

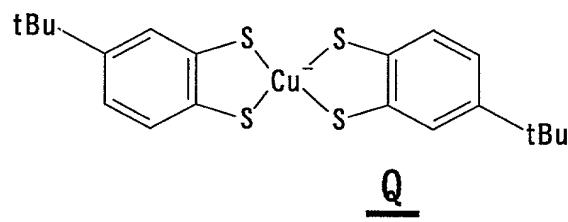
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A 1



A 2



As an organic solvent, diacetonealcohol was used. And, the content of the dye in a coating solution was 5% by weight. And, the thickness of the recording layer was 200 nm.

Subsequently, for each disk, a reflection layer was formed by a sputtering method. The thickness of the reflection layer and the composition thereof were as follows:

Reflection layer (80 nm): Au

Then, a protecting film (6) was formed on the reflection layer (5). The protecting film (6) was formed by coating an ultraviolet ray-curable resin: SD-211 (manufactured by Dainippon Ink & Chemicals. Inc.) with a spin coating method and performing ultraviolet ray-curing. The thickness of the protecting film after curing was 5  $\mu\text{m}$ .

Then, an ink receiving layer (7) was formed on the protecting film (6) of this disk (I) as follows:

Each component of the following composition was stirred for 1 hour in a mixer to prepare a coating solution No. 11.

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(Composition of the coating solution No. 11)

Monofunctional monomer:

a-1 (Aronix M-5600, manufactured by Toagoseikagaku (K.K.))

30 parts by weight

Acryloylmorpholine

10 parts by weight

Multifunctional monomer:

b-1 (A-600, manufactured by Shin-nakamura Kagakukogyo (K.K.))

20 parts by weight

Polymerization initiator:

Irgacure 651 (manufactured by Chiba Geigy)

5 parts by weight

Organic filler:

Staphyloid AC-3832 (particle size, 0.3  $\mu\text{m}$ )

30 parts by weight

Inorganic filler:

Silcron G-600 (particle size, 4.7  $\mu\text{m}$ ) 5 parts by weight

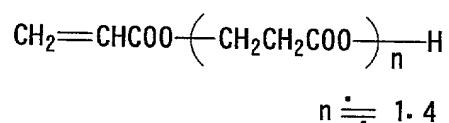
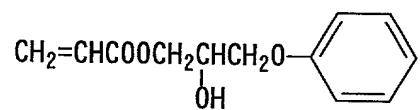
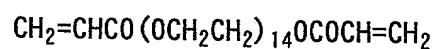
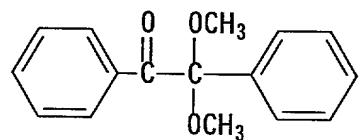
By using this coating solution No. 11, a coated film was formed on the protecting film (6) of the aforementioned disk using a screen printing method. This coated film was irradiated with an ultraviolet ray using an extra-high pressure mercury lamp so that the lamp intensity on the irradiated surface was 500  $\text{mJ}/\text{cm}^2$  to cure, to form an ink receiving layer (7) of about 10  $\mu\text{m}$ . The optical recording medium thus obtained is called sample No. 11.

Coating solution No. 12 was prepared according to the same manner as No. 11 except that trimethylolpropane triacrylate was used instead of monomer b-1 in the coating solution No. 11 and this coating solution No. 12 was used to form an ink receiving layer similarly. This optical recording medium is called sample No. 12.

Coating solution No. 13 was prepared according to the same manner as No. 11 except that 25 parts by weight of monomer a-2 (Aronix M-5700, manufactured by Toagoseikagaku (K.K.)) was used instead of 30 parts by weight of monomer a-1 and 15 parts by weight of polyethylene glycol 90 methacrylate was used instead of 10 parts by weight of acryloylmorpholine in the coating solution No. 11, respectively, and this coating solution No. 13 was used to form an ink receiving layer similarly. This optical recording medium is called sample No. 13.

Coating solution No. 14 was prepared according to the same manner as No. 13 except that dimethylaminopropylacrylamide was used instead of monomer a-2 (Aronix M-5700) in the coating solution No. 13 and this coating solution No. 14 was used to form an ink receiving layer similarly. This optical recording medium is called sample No. 14.

The structures of Aronix M-5600, Aronix M-5700, A-600, and Irgacure 651 used in this Example are shown below:

**Aronix M-5600****Aronix M-5700****A-600****Irgacure-651**

## [Comparative Example 1]

Coating solutions No. 15 and No. 16 were prepared, respectively, according to the same manner as in Example 1 except that the organic filler: Staphyloid AC-3832 was not added and the inorganic filler: Silcron G-600 was added in an amount of 35 parts by weight in the coating solutions No. 11 and No. 12 in Example 1 and the coating solutions No. 15 and No. 16 were used, respectively, to form an ink receiving layer similarly. These optical recording media are called samples No. 15 and No. 16.

Coating solutions No. 17 and No. 18 were prepared, respectively, according to the same manner as in Example 1 except that the inorganic filler: Silcron G-600 was not added and the organic filler: Staphyloid AC-3832 was added in an amount of 35 parts by weight in the coating solutions No. 13 and No. 14 in Example 1 and the coating solutions No. 17 and No. 18 were used, respectively, to form an ink receiving layer similarly. These optical recording media are called samples No. 17 and No. 18.

On the ink receiving layer surfaces of the resulting samples No. 11 to No. 18, printing was performed using an ink jet color printer (manufactured by Mastermind, myPRINTLAB). As an ink, color ink for Canon BJC600J was used.

For printability, surface property, resolution degree

and drying property were evaluated as follows:

(Surface property)

The surfaces after screen printing were observed and evaluated as follows:

- ◎ : Very smooth and dry on the surface
- : Smooth and no sticking feeling
- △ : Smooth on the surface but sticking feeling when touched with fingers
- × : Remaining mesh trace and bubble trace on the surface

(Resolution degree)

At an overlapping portion between ink and ink or a boundary portion of printing:

- ◎ : No blot
- : Blot of 0.4 mm or smaller
- △ : Blot of larger than 0.4 mm and smaller than 0.8 mm
- × : Blot of larger than 0.8 mm and repellence of ink

(Drying property)

Printing was performed and, after printing, time until an ink becomes unadhesive to fingers when contact with the printed surface is adopted as a drying time and evaluation was performed according to the following criteria:

- ◎ : Drying time is 30 seconds or shorter.
- : Drying time is longer than 30 seconds and 2 minutes or

shorter.

△ : Drying time is longer than 2 minutes and 10 minutes or shorter.

× : Drying time is longer than 10 minutes.

As regards water-resistance, printing was performed and, after drying the printing, the printed surface was dipped in flowing water for 1 minute and ink adsorbability and film property were evaluated as follows:

(Ink adsorbability)

After and before the printed surface was dipped in flowing water for 1 minute, the printed characters and images were observed with naked eyes and the change in the concentration was investigated.

○ : Little change in the concentration of printing

△ : The concentration is slightly lower but the printing characters are readable.

× : The printing was lost and becomes unreadable.

(Film property)

After and before the printed surface was dipped in flowing water for 1 minute, the state of the coating layer was observed with naked eyes.

○ : Little change in the layer surface

△ : The layer surface is slightly softened.

× : The layer flows with flowing water or peeled.

The results of the above performance evaluation on each optical recording medium sample are shown in Table 1. From Table 1, it is clear that the ink receiving layers of the samples No. 11 to No. 14 in accordance with the present invention have practically sufficient performance in an ink jet recording method. On the other hand, the ink receiving layers of the comparative samples No. 15 and No. 16 are excellent in surface property but inferior in resolution degree, drying property and ink adsorbability. In addition, the ink receiving layers of the comparative samples No. 17 and No. 18 are inferior in surface property and film property.

Table 1

Sample	<Printability>			<Water-Resistance>	
	Surface Property	Resolution Degree	Drying Property	Ink Adsorbability	Film Property
<b>Example 1</b>					
No. 11	○	○	○	○	○
No. 12	○	○	○	○	○
No. 13	○	◎	◎	○	○
No. 14	○	◎	◎	○	○
<b>Comparative Example 1</b>					
No. 15	◎	△	×	△	○
No. 16	◎	△	×	△	○
No. 17	×	○	○	○	△
No. 18	×	○	○	○	△

## [Example 2 and Comparative Example 2]

Coating solutions No. 21 to No. 28 were prepared, respectively, according to the same manner as in Example 1 and Comparative Example 1 except that Flowbeads EA209 (particle size, 10  $\mu\text{m}$ ) was used instead of the organic filler: Staphyloid AC-3832 and Aerosil TT-600 (particle size, 0.04  $\mu\text{m}$ ) was used instead of the inorganic filler: Silcron G-600 in the respective coating solutions No. 11 to No. 18 used in Example 1 and Comparative Example 1 and these respective coating solutions No. 21 to No. 28 were used to form an ink receiving layer similarly. These respective optical recording media are called samples No. 21 to No. 28.

For the resulting samples No. 21 to No. 28, the same evaluation as that in Example 1 was performed. As shown in Table 2, it is clear that the ink receiving layers of the samples No. 21 to No. 24 in accordance with the present invention have practically sufficient performance in an ink jet recording method. On the other hand, the ink receiving layers of the comparative samples No. 25 and No. 26 are excellent in surface property but inferior in resolution degree. In addition, the ink receiving layer of the comparative samples No. 27 and No. 28 are inferior in surface property and film property.

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Table 2

Sample	<Printability>			<Water-Resistance>	
	Surface Property	Resolution Degree	Drying Property	Ink Adsorbability	Film Property
<b>Example 2</b>					
No. 21	○	○	○	○	○
No. 22	○	○	○	○	○
No. 23	○	◎	◎	○	○
No. 24	○	◎	◎	○	○
<b>Comparative Example 2</b>					
No. 25	◎	△	△	△	○
No. 26	◎	△	△	△	○
No. 27	×	○	○	○	×
No. 28	×	○	○	○	×

## [Example 3 and Comparative Example 3]

Coating solutions No. 31 to No. 34 and No. 37 to No. 38 were prepared, respectively, according to the same manner as in Example 2 and Comparative Example 2 except that an animal protein fine particle (particle size, 5  $\mu\text{m}$ ) was used instead of the organic filler: Flowbeads EA209 in the respective coating solutions No. 21 to No. 24, and No. 27 to No. 28 used in Example 2 and Comparative Example 2 and these respective coating solutions were used to form an ink receiving layer similarly. These respective optical recording media are called samples No. 31 to No. 34, and No. 37 to No. 38.

For the resulting samples No. 31 to No. 34, and No. 37 and No. 38, the same evaluation as that in Example 1 was performed.

As shown in Table 3, it is clear that the ink receiving layers of the samples No. 31 to No. 34 in accordance with the present invention have practically sufficient performance in an ink jet recording method. On the other hand, the ink receiving layers of the comparative samples No. 37 and No. 38 are inferior in surface property and film property.

Table 3

Sample	<Printability>			<Water-Resistance>	
	Surface Property	Resolution Degree	Drying Property	Ink Adsorbability	Film Property
<b>Example 3</b>					
No. 31	○	○	○	○	○
No. 32	○	○	○	○	○
No. 33	○	◎	◎	○	○
No. 34	○	◎	◎	○	○
<b>Comparative Example 3</b>					
No. 37	×	○	○	○	△
No. 38	×	○	○	○	△

## [Example 4]

This Example is an embodiment in which a disk (II) having a protecting film (6) composed of two layers was used instead of the disk (I) having the protecting film (6) composed of one layer.

According to the same manner as the formation of the disk (I) in Example 1, the formation up to a reflection layer (5)

was performed. On this reflection layer (5), an ultraviolet curable-type resin: SD-211 was coated by a spin coating method, which was ultraviolet-cured to form a lower protecting film having a thickness of 5  $\mu\text{m}$  after curing. Subsequently, on the lower protecting film, an upper protecting film having a thickness of 8  $\mu\text{m}$  was formed using a white ink for screen printing (SSD F27 WHITE) manufactured by Dainippon Ink & Chemicals. Ink. to make a disk (II).

Then, on the upper protecting film of this disk (II), an ink receiving layer (7) was formed using each of the coating solutions No. 11 to No. 14 used in Example 1 similarly. These respective optical recording media are called samples No. 41 to No. 44.

In addition, the coating solutions No. 21 to No. 24 used in Example 2 were used, respectively, to form each ink receiving layer (7) similarly. These respective optical recording media are called samples No. 51 to No. 54.

Further, the coating solutions No. 31 to No. 34 used in Example 3 were used, respectively, to form each ink receiving layer (7) similarly. These respective optical recording media are called samples No. 61 to No. 64.

For the resulting samples No. 41 to No. 44, No. 51 to No. 54, and No. 61 to No. 64, the same evaluation as that in Example 1 was performed. As shown in Table 4, it is clear that the ink receiving layers of these samples have practically sufficient

performance in an ink jet recording method. Since these samples contained a white pigment in the upper protecting film, they had good and clear color development.

Table 4

Sample	<Printability>			<Water-Resistance>	
	Surface Property	Resolution Degree	Drying Property	Ink Adsorbability	Film Property
<b>Example 4</b>					
No. 41	○	○	○	○	○
No. 42	○	○	○	○	○
No. 43	○	◎	◎	○	○
No. 44	○	◎	◎	○	○
No. 51	○	○	○	○	○
No. 52	○	○	○	○	○
No. 53	○	◎	◎	○	○
No. 54	○	◎	◎	○	○
No. 61	○	○	○	○	○
No. 62	○	○	○	○	○
No. 63	○	◎	◎	○	○
No. 64	○	◎	◎	○	○

The present invention can be carried out in other various embodiments without departing from the spirit or main features thereof. For that reason, the foregoing Examples are merely illustration in all respects and are not construed to be limiting. Further, any changes belonging to the equivalent scope of claims are all within the scope of the present

invention.

## INDUSTRIAL APPLICABILITY

According to the coating composition of the present invention, as described above, by radiation-curing this composition, there can be formed an ink receiving layer having image clearness in an ink jet recording method, water-resistance and adhesion with substrates.

Therefore, the coating composition of the present invention is very useful for forming an ink receiving layer of the conventional ink jet recording material and, particularly, an ink receiving layer of an optical recording medium.

According to the optical recording medium of the present invention, since the medium has an ink receiving layer formed of this coating composition, that is provided a medium to be recorded thereon having the surface layer to be recorded thereon which is excellent in color printing compared with the prior art, has no ink blot or repellence or sticking. In particular, since the ink receiving layer is excellent in quick-drying, it is suitable in printing of high resolution graphics and photographs, and excellent printing having no color blot can be performed. Further, even when wiped after dipped in flowing water, printing fall and whole layer separation are not caused, leading to excellent water-resistance.

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## CLAIMS

1. A composition for coating which comprises a radiation-curable compound (A) containing a monofunctional (meth) acrylic monomer and a multifunctional more than bifunctional acrylic monomer, a water-absorbing organic filler (C) and a water-absorbing inorganic filler (D).

2. The composition for coating according to claim 1, wherein the radiation-curable compound (A) contains 50 to 90% by weight of the monofunctional (meth) acrylic monomer and 10 to 50% by weight of the multifunctional more than bifunctional (meth) acrylic monomer.

3. The composition for coating according to claim 2, which contains 10 to 60% by weight of the water-absorbing organic filler (C) and 1 to 20% by weight of the water-absorbing inorganic filler (D) relative to the radiation-curable compound (A).

4. The composition for coating according to claim 3, a particle size of the fillers (C) and (D) is 0.001 to 20  $\mu\text{m}$ .

5. The composition for coating according to claim 4, which contains 0.001 to 10% by weight of a polymerization initiator relative to the radiation-curable compound (A).

6. A recording material comprising an ink receiving layer composed of a radiation-cured composition for coating according to any one of claims 1 to 5 on at least one side of a hydrophobic supporting substrate.

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7. An optical recording medium which comprises an ink receiving layer composed of a radiation-cured composition for coating according to any one of claims 1 to 5 on the surface of the optical recording medium opposite to the surface on which optical writing/reading is performed.

## ABSTRACT

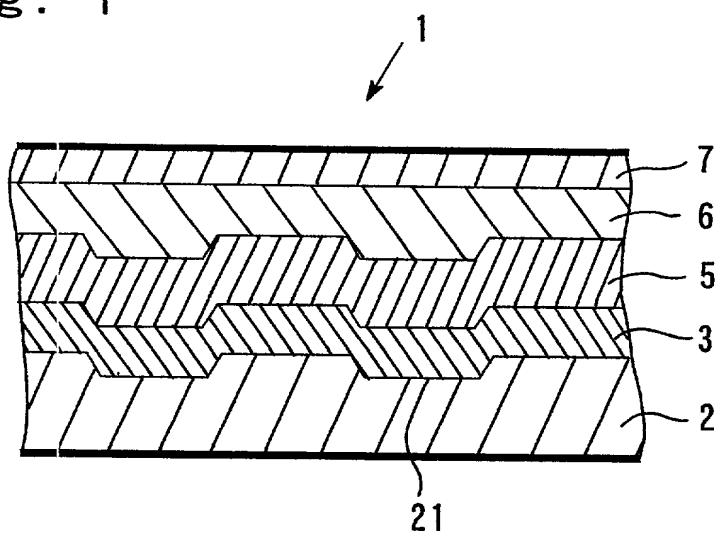
There are provided a composition for coating which can form an ink receiving layer having excellent adhesion with hydrophobic substrates, water-resistance and printing stability, and an optical recording medium having an ink receiving layer formed from this composition.

The composition contains a radiation-curable compound (A) composed of 50 to 90% by weight of a monofunctional (meth) acrylic monomer and 10 to 50% by weight of a multifunctional (meth) acrylic monomer, a water-absorbing organic filler (C) in an amount of 10 to 60% by weight relative to (A), and a water-absorbing inorganic filler (D) in an amount of 1 to 20% by weight relative to (A). This composition is applied on a protecting film (6) and cured to form an ink receiving layer (7), to obtain an optical disk (1).

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Fig. 1



**Declaration For U.S. Patent Application**

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled  
(Insert Title) COATING COMPOSITION AND OPTICAL RECORDING MEDIUM

the specification of which is attached hereto unless the following box is checked:

was filed on September 25, 1998 as PCT International Application  
 Number PCT/JP98/04286 and was amended on \_\_\_\_\_  
 and/or was filed on \_\_\_\_\_ as United States Application  
 Number \_\_\_\_\_ and was amended on \_\_\_\_\_.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claim(s), as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 C.F.R. §1.56.

I hereby claim foreign priority benefits under 35 U.S.C. §119(a)-(d) or §365(b) of any foreign application(s) for patent or inventor's certificate, or §365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below any foreign application for patent or inventor's certificate or PCT International Application having a filing date before that of the application(s) for which priority is claimed:

(List prior foreign applications. See note A on back of this page)	<u>Pat. 9-279651</u> (Number)	<u>Japan</u> (Country)	<u>27/September/1997</u> (Day/Month/Year Filed)	Priority Claimed
	<u>(Number)</u>	<u>(Country)</u>	<u>(Day/Month/Year Filed)</u>	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
	<u>(Number)</u>	<u>(Country)</u>	<u>(Day/Month/Year Filed)</u>	<input type="checkbox"/> Yes <input type="checkbox"/> No

I hereby claim the benefit under 35 U.S.C. §119(e) of any United States provisional application(s) listed below.

(Application Number) (Filing Date)(Application Number) (Filing Date)

(See Note B on back of this page)

 See attached list for additional prior foreign or provisional applications.

I hereby claim the benefit under 35 U.S.C. §120 of any United States application(s) or §365(c) of any PCT International application(s) designating the United States of America listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior application(s) (U.S. or PCT) in the manner provided by the first paragraph of 35, U.S.C. §112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 C.F.R. §1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

(List prior U.S. Applications or PCT International applications designating the U.S.)	<u>(Application Serial No.)</u>	<u>(Filing Date)</u>	<u>(Status) (patented, pending, abandoned)</u>
	<u>(Application Serial No.)</u>	<u>(Filing Date)</u>	<u>(Status) (patented, pending, abandoned)</u>

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further, that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

(See Note C on back of this page)

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February 23, 2000

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